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1386/10

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Examiner Initials*	Cite No.	Foreign Patent Document	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T <sup>6</sup>
		Country Code <sup>2</sup> - Number <sup>4</sup> - Kind Code <sup>3</sup> (if known)				
	3	WO 2005/014863	02-17-2005	Bionomics Limited		
	4	WO 2004/085674	10-07-2004	Bionomics Limited		
	5	WO 2002/050096	06-27-2002	Bionomics Limited		
	6	WO 2002/006521	01-24-2002	Bionomics Limited		

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Examiner Initials*	Cite No.	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.		T <sup>2</sup>
	7	Chou et al., "The lack of association between febrile convulsions and polymorphisms in SCN1A," <i>Epilepsy Research</i> , Vol. 54, pgs. 53-57 (2003).		
	8	Fujiwara et al., "Mutations of sodium channel a subunit type 1 (SCN1A) in intractable childhood epilepsies with frequent generalized tonic-clonic seizures," <i>Brain</i> , Vol. 126, pgs. 531-546 (2003).		
	9	Hirschhorn et al., "A comprehensive review of genetic association studies," <i>Genetics in Medicine</i> , Vol. 4, No. 2, pgs. 45-61 (2002).		
	10	Notification Concerning Transmittal of Copy of International Preliminary Report on Patentability for International Application No. PCT/AU2006/000841 dated January 3, 2008.		
	11	Official Action for U.S. Patent Application Serial No. 10/482,834 dated August 2, 2007.		
	12	Official Action for U.S. Patent Application Serial No. 10/482,834 dated April 4, 2008.		
	13	Official Action for U.S. Patent Application Serial No. 11/262,647 dated February 15, 2008.		
	14	Ohmori et al., "Significant correlation of the SCN1A mutations and severe myoclonic epilepsy in infancy," <i>Biochemical and Biophysical Research Communications</i> , Vol. 295, pgs. 17-23 (2002).		
	15	Staflstrom et al., "Epilepsy Genes: The Link Between Molecular Dysfunction and Pathophysiology," <i>Mental Retardation and Developmental Disabilities Research Reviews</i> , Vol. 6, pgs. 281-292 (2000).		
	16	Supplementary European Search Report corresponding to Australian Patent No. AU0200910 dated February 17, 2005.		

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